AMENDMENTS TO THE CLAIMS:

LISTING OF CLAIMS:

- 1. (Withdrawn) A method for regulating a menstrual cycle comprising administering a selective progesterone receptor modulator during a first dosing period and at least one progesterone during
- 2. (Withdrawn) The method of claim 1 wherein the first dosing period is between about 1 month and about 12 months.
- 3. (Withdrawn) The method of claim 2 wherein the second dosing period is between 1 day and 31 days.
- 4. (Withdrawn) The method of claim 3 wherein the second dosing period begins the first day after the first dosing period ends.
- 5. (Withdrawn) The method of claim 1 wherein the first dosing period and second dosing period overlap for at least one day.
- 6. (Withdrawn) The method of claim 1 wherein the SPRM is administered in an amount between 0.125 mg and 100 mg per day during the first dosing period.
- 7. (Withdrawn) The method of claim 6 wherein the progestogen is administered in an amount between 0.01 mg and 100 mg per day during the second dosing period.

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8. (Withdrawn) The method of claim 1 wherein the SPRM is selected form the group consisting of 11β [4-(hydroxyiminomethyl)phenyl]-17 β -methoxy-17- α -(methoxymethyl)estra-4,9-dien-3-one (asoprisnil), 11 β -4-(hydroxyiminomethyl)phenyl]-17 β -hydroxy-17 α .-(methoxymethyl)estra-4,9-dien-3-one (J912), and 11 β -[4-[(ethylaminocarbonyl-)oximinomethyl]phenyl]-

17 β -methoxy-17 α .-(methoxymethyl)estra-4,9- -dien-3-one (J956) as well as pharmaceutically acceptable salts thereof.

- 9. (Withdrawn) The method of claim 1 wherein the progestogen is selected from the group consisting of medroxyprogesterone, cyproterone, drospirenone, dydrogesterone, dienogest, noresthisterone, levonorgestrel, gestodene, promegestone, trimegestone, and pharmaceutically acceptable salts thereof.
- 10. (Withdrawn) The method of claim 9 wherein the method further comprises administering an estrogen during the second dosing period.
- 11. (Currently Amended) A method of treating a gynaecological disorder, the method comprising the steps of:

administering to a patient a SPRM for a first dosing period to achieve a therapeutic effect; and

administering at least one progestogen during a second dosing period to induce a predictable return to menstruation,

wherein the gynaecological disorder is <u>selected from the group consisting of uterine</u> fibroids, endometriosis, hormone replacement therapy, menorrhagia, metorrhagia, dysmenorrhea, adenomyosis or peritoneal adhesions.

- 12. (Original) The method of claim 11 wherein the first dosing period is between about 1 month and 12 months and the second dosing period is between 1 day and 31 days and the second dosing period begins the day following the first dosing period.
- 13. (Original) The method of claim 12 wherein the SPRM is selected from the group consisting of 11β [4-(hydroxyiminomethyl)phenyl]-17 β -methoxy-17-x-(methoxymethyl)estra-4,9-dien-3-one (asoprisnil), 11 β -[4-(hydroxyiminomethyk)phenyl]-17 β -hydroxy-17x-mrtho-xymethyl)estra-4,9-dien-3-one(J912), and 11 β -[4-[(ethylaminocarbonyl-)oximinomethyl]phenyl]-17 β -methoxy-17 α -(methoxymethyl)estra-4,9-dien-3-one (J956) as well as pharmaceutically acceptable salts thereof.

- 14. (Withdrawn) A kit comprising a SPRM and at least one progestogen.
- 15. (Withdrawn) The kit of claim 14 wherein the SPRM is selected form the group consisting of 11 β -[4-(hydroxyiminomethyl)phenyl]-17 β -methoxy-17- α -(methoxymethyl)estra-4,9-dien-3-one (asoprisnil), 11 β –[4-(hydroxyiminomethyl)phenyl]-17 β -hydroxy-17 α -(methoxymethyl)estra-4,9-dien-3-one (J912), and 11 β -[4-[(ethylaminocarbonyl-)oximinomethyl]phenyl]-17 β -methoxy-17 α –(methoxymethyl)estra-4,9--dien-3-one (J956) as well as pharmaceutically acceptable salts thereof; and the progestogens are selected from the group consisting of progesterone and any other synthetic progestin as well as their pharmaceutically acceptable salts and combinations of the foregoing.